



beta-propeller protein-associated neurodegeneration

Beta-propeller protein-associated neurodegeneration (BPAN) is a disorder that damages the nervous system and is progressive, which means that it gradually gets worse. Affected individuals develop a buildup of iron in the brain that can be seen with medical imaging. For this reason, BPAN is classified as a type of disorder called neurodegeneration with brain iron accumulation (NBIA), although the iron accumulation may not occur until late in the disease.

Many people with BPAN have recurrent seizures (epilepsy) beginning in infancy or early childhood. Several different types of seizures can occur in this disorder, even in the same individual. Often the first type to occur are febrile seizures, which are triggered by a high fever. Affected individuals can also experience generalized tonic-clonic seizures (also known as grand mal seizures). This type of seizure affects the entire body, causing muscle rigidity, convulsions, and loss of consciousness. Other seizure types that can occur in this disorder include short lapses in awareness that can have the appearance of staring spells or daydreaming (absence seizures, also called petit mal seizures), sudden episodes of weak muscle tone (atonic seizures), involuntary muscle twitches (myoclonic seizures), or more pronounced movements called epileptic spasms. Some individuals have seizure patterns that resemble those in epileptic syndromes, such as West syndrome or Lennox-Gastaut syndrome.

Children with BPAN also have intellectual disability, delayed development including significant problems with vocabulary and producing speech (expressive language), and difficulty coordinating movements (ataxia). Ataxia can affect the ability to walk and perform fine motor skills such as using utensils. Affected individuals can have behavioral changes that are often compared to features of a disorder called Rett syndrome. These features include repeated hand wringing or claspings (stereotypic hand movements); teeth grinding (bruxism); sleep disturbances; and problems with communication and social interaction characteristic of autism spectrum disorder.

In late adolescence or early adulthood, individuals with BPAN may begin to experience a gradual loss of intellectual functioning (cognitive decline) that can lead to a severe loss of thinking and reasoning abilities (dementia). Worsening problems with movement also occur, including dystonia and parkinsonism. Dystonia is a condition characterized by involuntary, sustained muscle contractions. In BPAN, the dystonia often starts in the arms. Parkinsonism can include unusually slow movement (bradykinesia), rigidity, tremors, an inability to hold the body upright and balanced (postural instability), and a shuffling walk that can cause recurrent falls.

The lifespan of people with BPAN varies. With proper management of their signs and symptoms, affected individuals can live into middle age. Death may result from

complications of dementia or movement problems, such as injuries from falls or swallowing difficulties (dysphagia) that can lead to a bacterial lung infection called aspiration pneumonia.

Frequency

BPAN is a rare disorder. Its prevalence is unknown, but it is thought to account for between 1 and 2 percent of all cases of NBIA disorders. About 100 affected individuals have been described in the medical literature. Some individuals who have been diagnosed with intellectual disability or early-onset parkinsonism based on their signs and symptoms have later been found to have BPAN when genetic testing was done.

Genetic Changes

BPAN is caused by mutations in the *WDR45* gene. This gene provides instructions for making the WIPI4 protein. WIPI4 has a characteristic structure resembling a seven-bladed propeller, from which the name of the disorder is derived. The WIPI4 protein is involved in a process called autophagy, which helps clear unneeded materials from cells, including excess amounts of an iron storage protein called ferritin.

Most of the *WDR45* gene mutations identified in people with BPAN result in the absence of functioning WIPI4 protein. As a result, the process of autophagy is impaired, making cells less efficient at removing damaged cell structures and waste materials. Researchers suggest that nerve cells (neurons) may be particularly vulnerable to impaired autophagy because they have long extensions (axons and dendrites), making it even more difficult to transport the waste materials from these structures to the compartments in the cell body where recycling takes place (the lysosomes). The waste materials can build up in these areas and damage them. Damage to neurons results in the neurological problems that occur in BPAN.

Inheritance Pattern

This condition is inherited in an X-linked dominant pattern. The gene associated with this condition is located on the X chromosome, which is one of the two sex chromosomes. In females (who have two X chromosomes), a mutation in one of the two copies of the gene in each cell is sufficient to cause the disorder. In males (who have only one X chromosome), a mutation in the only copy of the gene in each cell causes the disorder. A characteristic of X-linked inheritance is that fathers cannot pass X-linked traits to their sons.

In most X-linked dominant disorders, males experience more severe symptoms than females. While this is not always the case in BPAN, most individuals with the disorder are females, likely because a smaller number of affected males survive until birth.

Almost all cases of BPAN result from new mutations in the gene and occur in people with no history of the disorder in their family. Rarely, an affected person inherits the

mutation from a mildly affected mother. Among reported cases, males with BPAN and most females with BPAN have not had children.

Other Names for This Condition

- BPAN
- NBIA5
- neurodegeneration with brain iron accumulation 5
- SENDA
- static encephalopathy of childhood with neurodegeneration in adulthood

Diagnosis & Management

Genetic Testing

- Genetic Testing Registry: Beta-propeller protein-associated neurodegeneration
<https://www.ncbi.nlm.nih.gov/gtr/conditions/CN168656/>

Other Diagnosis and Management Resources

- GeneReview: Beta-Propeller Protein-Associated Neurodegeneration
<https://www.ncbi.nlm.nih.gov/books/NBK424403>

General Information from MedlinePlus

- Diagnostic Tests
<https://medlineplus.gov/diagnostictests.html>
- Drug Therapy
<https://medlineplus.gov/drugtherapy.html>
- Genetic Counseling
<https://medlineplus.gov/geneticcounseling.html>
- Palliative Care
<https://medlineplus.gov/palliativecare.html>
- Surgery and Rehabilitation
<https://medlineplus.gov/surgeryandrehabilitation.html>

Additional Information & Resources

MedlinePlus

- Encyclopedia: Intellectual Disability
<https://medlineplus.gov/ency/article/001523.htm>
- Encyclopedia: Movement -- Uncontrollable
<https://medlineplus.gov/ency/article/003201.htm>

- Encyclopedia: Movement -- Uncoordinated
<https://medlineplus.gov/ency/article/003198.htm>
- Encyclopedia: Neurodegeneration with Brain Iron Accumulation
<https://medlineplus.gov/ency/article/001225.htm>
- Health Topic: Degenerative Nerve Diseases
<https://medlineplus.gov/degenerativenervediseases.html>
- Health Topic: Dementia
<https://medlineplus.gov/dementia.html>
- Health Topic: Epilepsy
<https://medlineplus.gov/epilepsy.html>
- Health Topic: Movement Disorders
<https://medlineplus.gov/movementdisorders.html>

Genetic and Rare Diseases Information Center

- Beta-Propeller Protein-Associated Neurodegeneration
<https://rarediseases.info.nih.gov/diseases/12570/beta-propeller-protein-associated-neurodegeneration>

Additional NIH Resources

- National Institute of Neurological Disorders and Stroke: Dystonias Information Page
<https://www.ninds.nih.gov/Disorders/All-Disorders/Dystonias-Information-Page>
- National Institute of Neurological Disorders and Stroke: Epilepsy Information Page
<https://www.ninds.nih.gov/Disorders/All-Disorders/Epilepsy-Information-Page>
- National Institute of Neurological Disorders and Stroke: Neurodegeneration with Brain Iron Accumulation
<https://www.ninds.nih.gov/Disorders/All-Disorders/Neurodegeneration-Brain-Iron-Accumulation-Information-Page>

Educational Resources

- Centers for Disease Control and Prevention: Developmental Disabilities
<https://www.cdc.gov/ncbddd/developmentaldisabilities/>
- Merck Manual Consumer Version: Dystonia
<http://www.merckmanuals.com/home/brain-spinal-cord-and-nerve-disorders/movement-disorders/dystonia>

- Merck Manual Consumer Version: Parkinsonism
<http://www.merckmanuals.com/home/brain-spinal-cord-and-nerve-disorders/movement-disorders/parkinsonism>
- Orphanet: Beta-propeller protein-associated neurodegeneration
http://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=329284

Patient Support and Advocacy Resources

- NBIA Disorders Association
<http://www.nbiadisorders.org/about-nbia/bpan>
- NBIAcure
<http://nbiacure.org/learn/nbia-disorders/bpan/>
- The Arc: for People with Intellectual and Developmental Disabilities
<http://www.thearc.org/>

GeneReviews

- Beta-Propeller Protein-Associated Neurodegeneration
<https://www.ncbi.nlm.nih.gov/books/NBK424403>

ClinicalTrials.gov

- ClinicalTrials.gov
<https://clinicaltrials.gov/ct2/results?term=%22beta-propeller+protein-associated+neurodegeneration%22+%5BDISEASE%5D+OR+NCT02587858+%5BID-NUMBER%5D>

Scientific Articles on PubMed

- PubMed
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28beta-propeller+protein-associated+neurodegeneration%5BTIAB%5D%29+AND+english%5BIa%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D>

OMIM

- NEURODEGENERATION WITH BRAIN IRON ACCUMULATION 5
<http://omim.org/entry/300894>

MedGen

- Beta-propeller protein-associated neurodegeneration
<https://www.ncbi.nlm.nih.gov/medgen/489843>

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